BLUESTOP MAX- external analgesic gel
Clavel Corporation

Blue Stop Max Roll-On

Active Ingredients

Lidocaine HCl 4%

Menthol 1%

Purpose

Topical Analgesic

Use

Temporary relief of pain

Warnings

For external use only

When using this product avoid contact with eyes

Stop use and ask a doctor if condition worsens or symptoms persist for more than 7 days or clear up and occur again within a few days.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Adults and children 2 years of age and older: Apply to affected area not more than 3 to 4 times daily.
- Children under 2 years of age: Consult a doctor.

Other Information

Store at room temperature

Avoid excessive hrat

Inactive Ingredients

Aloe Vera Leaf, Alpha-Tocopherol Acetate, Ammonium Acryloyldimethyltaurate, Dimethylacrylamide, Lauryl Methacrylate and Laureth-4 Methacrylate Copolymer, Trimethylolpropane Triacrylate Crosslinked (45000 Mpa.S), Cetyl Alcohol, Coconut Oil,

Dimethyl Sulfone, Ethylhexylglycerin,Ethylhexyl Palmitate, Fragrance, Garcinia Indica Seed Butter, Glucosamine Sulfate, Glycerin, Melaleuca Alternifolia Leaf, Phenoxyethanol, Sodium Acrylate/Sodium Acryloyldimethyltaurate Copolymer (4000000 MW), Vitamin A Palmitate, Water, Witch Hazel

Questions or Comments

1-800-432-5464

Roll-On Label and Carton Label

BLUESTOP MAX

external analgesic gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82714-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	10 mg in 1 g
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	40 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
WITCH HAZEL (UNII: 101I4J0U34)	
MELALEUCA ALTERNIFOLIA LEAF (UNII: G43C57162K)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
WATER (UNII: 059QF0KO0R)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SODIUM ACRYLATE/SODIUM ACRYLOYLDIMETHYLTAURATE COPOLYMER (4000000 MW) (UNII: 1DXE3F3OZX)	
GARCINIA INDICA SEED BUTTER (UNII: US2H3D7800)	
GLUCOSAMINE SULFATE (UNII: 1FW7WLR731)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
COCONUT OIL (UNII: Q9L0O73W7L)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
ETHYLHEXYL PALMITATE (UNII: 2865993309)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE, DIMETHYLACRYLAMIDE, LAURYL METHACRYLATE AND LAURETH-4 METHACRYLATE COPOLYMER, TRIMETHYLOLPROPANE TRIACRYLATE CROSSLINKED (45000 MPA.S) (UNII: Q7UI015FF9)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82714-001-01	1 in 1 CARTON	01/10/2023	
1		85 g in 1 APPLICATOR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	01/10/2023	

Labeler - Clavel Corporation (927855643)

Establishment

Name	Address	ID/FEI	Business Operations
TAKA USA Inc, dba Cosmetic Innovations		802860515	manufacture(82714-001)

Revised: 1/2026 Clavel Corporation